

FILE 'HOME' ENTERED AT 18:45:43 ON 25 JUN 2010

=> file hcaplus  
COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
0.22	0.22

FULL ESTIMATED COST

FILE 'HCAPLUS' ENTERED AT 18:46:29 ON 25 JUN 2010  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.  
COPYRIGHT (C) 2010 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 25 Jun 2010 VOL 153 ISS 1  
FILE LAST UPDATED: 24 Jun 2010 (20100624/ED)  
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Apr 2010  
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Apr 2010

HCAPLUS now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2010.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate substance identification.

```
=> s (salmeterol (P) (dimerization or dimerisation))
      1788 SALMETEROL
      51734 DIMERIZATION
      91 DIMERISATION
L1      0 (SALMETEROL (P) (DIMERIZATION OR DIMERISATION))
```

```
=> s (salmeterol and (dimerization or dimerisation or decomposition or dimer or
degrad?))
```

```
      1788 SALMETEROL
      51734 DIMERIZATION
      91 DIMERISATION
      224560 DECOMPOSITION
      480474 DECOMPN
      581046 DECOMPOSITION
            (DECOMPOSITION OR DECOMPN)
      125090 DIMER
      342813 DEGRAD?
      391724 DEGRDN
      580856 DEGRAD?
            (DEGRAD? OR DEGRDN)
L2      25 (SALMETEROL AND (DIMERIZATION OR DIMERISATION OR DECOMPOSITION
OR DIMER OR DEGRAD?))
```

=> s L2 and (acid adj (mineral or organic or inorganic or nitric of sulfuric or sulphuric or phosphoric))

MISSING OPERATOR 'ADJ (MINERAL'

The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> s L2 and (acid (W2) (mineral or organic or inorganic or nitric of sulfuric or sulphuric or phosphoric))

MISSING OPERATOR 'ACID (W2'

The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> s L2 and (acid (5A) (mineral or organic or inorganic or nitric of sulfuric or sulphuric or phosphoric))

5078847 ACID

424957 MINERAL

479407 ORGANIC

1209836 ORG

1347273 ORGANIC

(ORGANIC OR ORG)

158457 INORGANIC

342573 INORG

426564 INORGANIC

(INORGANIC OR INORG)

239715 NITRIC

187029 SULFURIC

1762 NITRIC OF SULFURIC

(NITRIC(1W) SULFURIC)

16312 SULPHURIC

127699 PHOSPHORIC

251369 ACID (5A) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC)

L3 1 L2 AND (ACID (5A) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC))

=> d L3 TI AB IBIB

L3 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2010 ACS on STN

TI A stable aerosol solution containing glucocorticoids suitable for oral or nasal inhalation

AB Aerosol soln. formulations contg. glucocorticosteroids stabilized by adding water or a mixt. of water and citric acid, avoiding corrosion of the elements of container under std. storage conditions are described. The formulations comprise 0.05 to 1.0% by wt. of a glucocorticoid having a C-20 ketone and OH group in carbons 17 and/or 21 as active substance; 0.10 to 3% by wt. of a selected stabilizer selected between water, or a mixt. of water and org. acid selected between citric acid and tartaric acid; a cosolvent in amt. sufficient to solubilize the active substance; optionally a surfactant; and propellant in sufficient amt. to achieve 100% by wt. of the finished soln. Glucocorticosteroids having a C-20 ketone and an OH group at the C-17 and/or 21 position with varying substituents, have many well-known therapeutic uses, esp. based upon their anti-inflammatory activity. This types of steroids, glucocorticosteroids, and their pharmaceutical formulations are useful in the treatment of several diseases including bronchial disorders and inflammatory conditions. Preferably, the glucocorticoid is selected between triamcinolone acetonide, budesonide, dexamethasone and betamethasone 17-valerate. A method for stabilizing aerosol pharmaceutical soln. formulations contg. glucocorticoids susceptible to oxidative degrdn. and use of a stabilizer selected between water and a mixt. of water and org.

acid selected between citric acid and tartaric acid are also described. For example, an aerosol compn. contg. 10 mL of a soln. of 150 mg of budesonide in 50 mL of ethanol and 174 mL water showed an increase in budesonide stability in presence of 0.333 g aluminum oxide compared to the compn. without addn. of water. The budesonide percentage found after 22 h of storage at 75.degree. were 14.7% and 4.9% for the aerosol formulations with and without water, resp.

ACCESSION NUMBER: 2005:393969 HCAPLUS  
DOCUMENT NUMBER: 142:417209  
TITLE: A stable aerosol solution containing glucocorticoids suitable for oral or nasal inhalation  
INVENTOR(S): Vega, Julio Cesar; De Bonis, Fabian  
PATENT ASSIGNEE(S): Laboratorio Pablo Cassara S.R.L., Argent.  
SOURCE: Eur. Pat. Appl., 13 pp.  
CODEN: EPXXDW  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1527772	A1	20050504	EP 2004-19514	20040817
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR				
IN 2004MU00888	A	20070525	IN 2004-MU888	20040816
IN 220652	A1	20080815		
BR 2004003316	A	20050621	BR 2004-3316	20040819
MX 2004008409	A	20050705	MX 2004-8409	20040830
US 20050095206	A1	20050505	US 2004-943403	20040917
PRIORITY APPLN. INFO.:			AR 2003-103969	A 20031030
OS.CITING REF COUNT:	1	THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)		
REFERENCE COUNT:	8	THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT		

=> d his

(FILE 'HOME' ENTERED AT 18:45:43 ON 25 JUN 2010)

FILE 'HCAPLUS' ENTERED AT 18:46:29 ON 25 JUN 2010

L1 0 S (SALMETEROL (P) (DIMERIZATION OR DIMERISATION))  
L2 25 S (SALMETEROL AND (DIMERIZATION OR DIMERISATION OR DECOMPOSITIO  
L3 1 S L2 AND (ACID (5A) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC

=> d que L2

L2 25 SEA FILE=HCAPLUS ABB=ON (SALMETEROL AND (DIMERIZATION OR DIMERISATION OR DECOMPOSITION OR DIMER OR DEGRAD?))

=> d que L3

L2 25 SEA FILE=HCAPLUS ABB=ON (SALMETEROL AND (DIMERIZATION OR DIMERISATION OR DECOMPOSITION OR DIMER OR DEGRAD?))  
L3 1 SEA FILE=HCAPLUS ABB=ON L2 AND (ACID (5A) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC))

=> s L2 and (acid (W) (mineral or organic or inorganic or nitric of sulfuric or sulphuric or phosphoric))

```

5078847 ACID
  424957 MINERAL
  479407 ORGANIC
1209836 ORG
1347273 ORGANIC
      (ORGANIC OR ORG)
  158457 INORGANIC
  342573 INORG
  426564 INORGANIC
      (INORGANIC OR INORG)
  239715 NITRIC
  187029 SULFURIC
    1762 NITRIC OF SULFURIC
      (NITRIC(1W)SULFURIC)
    16312 SULPHURIC
  127699 PHOSPHORIC
    8394 ACID (W) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC OF SULFURIC
      OR SULPHURIC OR PHOSPHORIC)
L4      0 L2 AND (ACID (W) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC OF
      SULFURIC OR SULPHURIC OR PHOSPHORIC))

```

```

=> s L2 and (acid (S) (citric or fumaric or ascorbic mineral or organic or
inorganic or nitric of sulfuric or sulphuric or phosphoric))

```

```

5078847 ACID
  113279 CITRIC
  26650 FUMARIC
  98756 ASCORBIC
  424957 MINERAL
    0 ASCORBIC MINERAL
      (ASCORBIC(W)MINERAL)
  479407 ORGANIC
1209836 ORG
1347273 ORGANIC
      (ORGANIC OR ORG)
  158457 INORGANIC
  342573 INORG
  426564 INORGANIC
      (INORGANIC OR INORG)
  239715 NITRIC
  187029 SULFURIC
    1762 NITRIC OF SULFURIC
      (NITRIC(1W)SULFURIC)
    16312 SULPHURIC
  127699 PHOSPHORIC
  419838 ACID (S) (CITRIC OR FUMARIC OR ASCORBIC MINERAL OR ORGANIC OR
      INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC)
L5      3 L2 AND (ACID (S) (CITRIC OR FUMARIC OR ASCORBIC MINERAL OR ORGAN
      IC OR INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC
      ))

```

```

=> S L5 NOT L2
L6      0 L5 NOT L2

```

```

=> d his

```

```

(FILE 'HOME' ENTERED AT 18:45:43 ON 25 JUN 2010)

```

```

FILE 'HCAPLUS' ENTERED AT 18:46:29 ON 25 JUN 2010

```

```

L1      0 S (SALMETEROL (P) (DIMERIZATION OR DIMERISATION))
L2      25 S (SALMETEROL AND (DIMERIZATION OR DIMERISATION OR DECOMPOSITIO
L3      1 S L2 AND (ACID (5A) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC

```

L4 0 S L2 AND (ACID (W) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC O  
 L5 3 S L2 AND (ACID (S) (CITRIC OR FUMARIC OR ASCORBIC MINERAL OR OR  
 L6 0 S L5 NOT L2

=> s L5 NOT L3  
 L7 2 L5 NOT L3

=> d L7 1-2 TI AB IBIB

L7 ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2010 ACS on STN  
 TI Nanoparticles and porous particles for aerosol inhalant formulations and  
 methods of making the same  
 AB The subject matter disclosed herein relates to compns. and methods for  
 engineering porous particles for aerosol formulations for pulmonary drug  
 delivery. Certain embodiments disclosed herein further relate to methods  
 for stabilizing suspension-based formulations in hydrofluoroalkane  
 propellants with nanoparticles. Thus, chitosan nanoparticles were prepd.  
 by std. nano-pptn. technique: chitosan was dissolved in water with the aid  
 of hydrochloric acid at 80.degree.C; hydrogen peroxide (30 %) was  
 subsequently added into the chitosan aq. soln. to degrade the long chain  
 chitosan to water-sol. smaller oligomers; the chitosan oligomer was then  
 dissolved in water and added into a large vol. of ethanol dropwise under  
 mild stirring; chitosan was then pptd. from the org. phase as nanoparticles  
 because it is insol. in ethanol. The surface of the chitosan nanoparticle  
 was then modified by ring-opening polymn. of the lactide with the hydroxy  
 and amine groups on chitosan nanoparticle surface as the initiator; the  
 nanoparticles have an av. size of approx. 100 nm, although the size can be  
 controlled by varying the prepn. parameters such as the concn. of  
 chitosan aq. soln., different type of org. solvents used and the  
 presence of surfactant.

ACCESSION NUMBER: 2010:627394 HCAPLUS  
 DOCUMENT NUMBER: 152:576514  
 TITLE: Nanoparticles and porous particles for aerosol  
 inhalant formulations and methods of making the same  
 INVENTOR(S): Da Rocha, Sandro Rp; Wu, Libo  
 PATENT ASSIGNEE(S): Wayne State University, USA  
 SOURCE: PCT Int. Appl., 38pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 2  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2010057214	A2	20100520	WO 2009-US64863	20091117
W:	AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PE, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, SE, SI, SK, SM, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
PRIORITY APPLN. INFO.:			US 2008-199518P	P 20081117
			US 2008-199519P	P 20081117

L7 ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2010 ACS on STN  
 TI Composition and method for topical treatment of tar-responsive dermatological disorders  
 AB The present invention relates to a compn. including a wax and a therapeutically effective amt. of tar for topical treatment of a tar-responsive dermatol. disorder, the compn. being in liq. or light gel form when at a temp. selected from room temp. and a temp. of skin of a mammal upon application of the compn. to the skin of the mammal. The invention also relates to a method of treating a tar-responsive dermatol. disorder by topically applying the compn. to skin of a mammal, preferably a human, that is affected by the disorder. Thus, a fast-drying liq. tar compn. was formulated contg. coal tar soln. 15 g, ethanol 42 g, propylene glycol 5 g, cyclomethicone (DC 345) 15 g, tri-Et citrate 5 g, Brij 93 10 g, liq. wax DIADD (dioctyldodecyl dodecanedioate) 5 g, and an optional fragrance 3 g. Topical application of the compn. for 4 mo to a human subject having plaque psoriasis resulted in 90% improvement of clin. signs of disorder.

ACCESSION NUMBER: 2007:993749 HCAPLUS  
 DOCUMENT NUMBER: 147:330433  
 TITLE: Composition and method for topical treatment of tar-responsive dermatological disorders  
 INVENTOR(S): Yu, Ruey J.; Van Scott, Eugene J.; Lee, Yaling  
 PATENT ASSIGNEE(S): Tristrata, Inc., USA  
 SOURCE: U.S. Pat. Appl. Publ., 15pp.  
 CODEN: USXXCO  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20070207222	A1	20070906	US 2007-680227	20070228
AU 2007223560	A1	20070913	AU 2007-223560	20070228
AU 2007223560	A2	20081016		
CA 2644311	A1	20070913	CA 2007-2644311	20070228
WO 2007103687	A2	20070913	WO 2007-US62975	20070228
WO 2007103687	A3	20081211		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA				
EP 1998788	A2	20081210	EP 2007-757636	20070228
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS				
JP 2009528382	T	20090806	JP 2008-557487	20070228
MX 2008011236	A	20090210	MX 2008-11236	20080902
CN 101460060	A	20090617	CN 2007-80015758	20081031
US 20100093827	A1	20100415	US 2009-638505	20091215
PRIORITY APPLN. INFO.:				
			US 2006-778128P	P 20060301
			US 2007-680227	A3 20070228
			WO 2007-US62975	W 20070228

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

=> d his

(FILE 'HOME' ENTERED AT 18:45:43 ON 25 JUN 2010)

FILE 'HCAPLUS' ENTERED AT 18:46:29 ON 25 JUN 2010

```
L1      0 S (SALMETEROL (P) (DIMERIZATION OR DIMERISATION))
L2      25 S (SALMETEROL AND (DIMERIZATION OR DIMERISATION OR DECOMPOSITIO
L3      1 S L2 AND (ACID (5A) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC
L4      0 S L2 AND (ACID (W) (MINERAL OR ORGANIC OR INORGANIC OR NITRIC O
L5      3 S L2 AND (ACID (S) (CITRIC OR FUMARIC OR ASCORBIC MINERAL OR OR
L6      0 S L5 NOT L2
L7      2 S L5 NOT L3
```

=> d que L2

```
L2      25 SEA FILE=HCAPLUS ABB=ON (SALMETEROL AND (DIMERIZATION OR
        DIMERISATION OR DECOMPOSITION OR DIMER OR DEGRAD?))
```

=> d que L3

```
L2      25 SEA FILE=HCAPLUS ABB=ON (SALMETEROL AND (DIMERIZATION OR
        DIMERISATION OR DECOMPOSITION OR DIMER OR DEGRAD?))
L3      1 SEA FILE=HCAPLUS ABB=ON L2 AND (ACID (5A) (MINERAL OR ORGANIC
        OR INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC))
```

=> d que L4

```
L2      25 SEA FILE=HCAPLUS ABB=ON (SALMETEROL AND (DIMERIZATION OR
        DIMERISATION OR DECOMPOSITION OR DIMER OR DEGRAD?))
L4      0 SEA FILE=HCAPLUS ABB=ON L2 AND (ACID (W) (MINERAL OR ORGANIC
        OR INORGANIC OR NITRIC OF SULFURIC OR SULPHURIC OR PHOSPHORIC))
```

=> d que L5

```
L2      25 SEA FILE=HCAPLUS ABB=ON (SALMETEROL AND (DIMERIZATION OR
        DIMERISATION OR DECOMPOSITION OR DIMER OR DEGRAD?))
L5      3 SEA FILE=HCAPLUS ABB=ON L2 AND (ACID (S) (CITRIC OR FUMARIC
        OR ASCORBIC MINERAL OR ORGANIC OR INORGANIC OR NITRIC OF
        SULFURIC OR SULPHURIC OR PHOSPHORIC))
```